

RESEARCH ARTICLE

Comparative analysis of the integument of different tree frog species from *Ololygon* and *Scinax* genera (Anura: Hylidae)

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ABSTRACT. The integuments of ten treefrog species of two genera from Scinaxnae – *O. angrensis* (Lutz, 1973), *O. flavoguttata* (Lutz & Lutz, 1939), *O. humilis* (Lutz & Lutz, 1954), *O. perpusilla* (Lutz & Lutz, 1939), *O. v-signata* (Lutz, 1968), *Scinax hayii* (Barbour, 1909), *S. similis* (Cochran, 1952), *O. trapicheroi* (Lutz & Lutz, 1954) and *S. x-signatus* (Spix, 1824) – were investigated using conventional and histochemical techniques of light microscopy, and polarized light microscopy. All integuments showed the basic structure of the anuran integument. Moreover, the secretory portions of exocrine glands, such as serous merocrine and apocrine glands, were found to be restricted to the spongy dermis. Lipid content occurred together with the heterogeneous secretory material of the glands with an apocrine secretion mechanism. In addition, clusters of these apocrine glands were present in the ventrolateral integument of some species. Melanophores were also visualized in all examined hylids. However, the occurrence of iridophores, detected through polarized light microscopy, varied according to the species. The Eberth-Katschenko layer occurred in the dorsal integument from both genera, but it was only present in the ventral integument of *O. albicans*, *O. angrensis*, *O. flavoguttata*, *O. perpusilla* and *O. v-signata*. Although the integument of all treefrogs showed the same basic structure, some characteristics were genus-specific; however, these features alone may not be used to distinguish both genera.

KEY WORDS. Brazilian Atlantic forest, histochemistry, hylids, treefrog.

INTRODUCTION

Many challenges confront biologists studying amphibians, since human activities have been prejudicial to natural biota. Hylidae is a large anuran family, and one of the most abundant and prominent groups of frogs in the Neotropics. Overall, 345 species of Hylidae are known to exist in Brazil (SBH 2016).

The integument of anurans performs several functions, such as protection against diverse environmental circumstances (Elkan 1968, Fox 1986a, b, Greven et al. 1995, de Brito-Gitirana and Azevedo 2005, Azevedo et al. 2006, de Brito-Gitirana and Azevedo 2005), mechanical protection (Fox 1986a, b, Azevedo et al. 2006), chemical defense (Delfino et al. 1995, Daly 1995, Jekel et al. 2015), sensory perception (Mearow and Diamond 1988, Koyama et al. 2001), ionic transport and water absorp-

tion (Sullivan et al. 2000, Azevedo et al. 2006), and respiration (Duellman and Trueb 1994).

In adult anurans, the integument consists of two firmly attached layers: the epidermis and the dermis, which is located just beneath the epidermis. The epidermis, formed by a stratified squamous epithelium, overlies the dermis of connective tissue, which is subdivided into two layers: the spongy dermis and the compact dermis. The spongy dermis is formed by loose connective tissue with pigment cells, such as melanophores (melanin producing cells) and iridophores (with reflective or iridescent structures) located just beneath the basal lamina (Bagnara et al. 1968). In addition to the pigment cells, blood vessels, alveolar mucous, serous, mixed and granular glands also occur. The compact dermis is composed of a series of alternating layers of collagenous fiber bundles arranged in a crisscross manner (Fox

1986a, b, Azevedo et al. 2005, de Brito-Gitirana and Azevedo 2005). Nevertheless, integument structure varies according to the body region (de Brito-Gitirana and Azevedo 2005, Felsemburgh et al. 2007). In addition, different types of exocrine glands occur in the anuran integument, with their secretory portions restricted to the spongy dermis (Duellmann and Trueb 1994, Brizzi et al. 2002).

In general, anurans show a wide array of colors related to specialized cells named chromatophores. Color and reflectivity are important mechanisms that allow anurans to change their integument color, enabling the maintenance of body temperature and avoiding detection by predators (Duellmann and Trueb 1994). Three types of chromatophores are important for amphibian coloration: melanophores (with a black or brownish pigment named melanin), xanthophores (with yellow colored pigments), and iridophores (with reflective or iridescent structures). These pigment cells are located in the spongy dermis just beneath the basal lamina, and their arrangement was called the Dermal Chromatophore Unit (DCU) (Bagnara et al. 1968). However, the exact mechanism of DCU organization is still unknown.

Some anurans, like arboreal frogs, exhibit diverse morphology in their typical integument outline, showing cutaneous adaptations to avoid evaporative water loss (Blaylock et al. 1976, Warburg et al. 2000, Barbeau and Lillywhite 2005). Such adaptations include exocrine glands secreting lipids, which are spread over the body surface, aided by an elaborate series of behavioral movements to form a useful barrier against water loss. In addition, the histological aspects of the anuran integument can contribute to the differentiation of anuran species from *Proceratophrys* Miranda-Ribeiro, 1920 (Felsemburgh et al. 2007).

The abundant fauna of amphibians of Brazil (SBH 2016) contrast with the present loss and degradation of natural habitats and the scarce knowledge on the morphological, taxonomical, biological, and geographical distribution of most species. The aim of the present work was to investigate the morphological features of the integuments of some hylids in order to characterize this tissue and determine whether there are species-specific structures, contributing to the knowledge of the integument biology.

MATERIAL AND METHODS

Nine species of male adult tree frogs belonging to *Scinax* from both *Oloolygon* Fitzinger, 1843 and *Scinax* Wagler, 1830 genera were collected in Serra dos Órgãos National Park (PARNASO, 22°29'31"S, 42°59'11.48"W), in the municipality of Teresópolis, during two years (from March 2007 to November 2008). *Oloolygon angrensis* (Lutz, 1973) was collected in Rio de Janeiro state at the Lídice district (22°46'53.82"S, 44°13'55.96"W), municipality of Rio Claro (from March 2009 to March 2010) (Table 1). Tree frogs were collected under license permit number 15396-1 and 219/2011, issued by the Instituto Chico Mendes de Conservação da Biodiversidade (ICMBio).

Samplings were carried out at night from 7 to 10 pm and all individuals collected (Figs 1–10) were carefully placed in

Table 1. List of individuals used in this study.

	Number of individuals	Common name (Frost 2016)
<i>Oloolygon albicans</i> (Bokermann, 1967)	10	Teresópolis Snouted Treefrog
<i>Oloolygon angrensis</i> (Lutz, 1973)	5	Serra da Bocaina Snouted Treefrog
<i>Oloolygon flavoguttata</i> (Lutz & Lutz, 1939)	5	Yellowbelly Snouted Treefrog
<i>Oloolygon humilis</i> (Lutz & Lutz, 1954)	3	Rio Babi Snouted Treefrog
<i>Oloolygon perpusilla</i> (Lutz & Lutz, 1939)	3	Bandeirantes Snouted Treefrog
<i>Oloolygon trapicheroi</i> (Lutz & Lutz, 1954)	5	Three-lined Snouted Treefrog
<i>Oloolygon v-signata</i> (Lutz, 1968)	2	Forest Snouted Treefrog
<i>Scinax similis</i> (Cochran, 1952)	3	Cochran's Snouted Treefrog
<i>Scinax hayii</i> (Barbour, 1909)	3	Hay's Snouted Treefrog
<i>Scinax x-signatus</i> (Spix, 1824)	3	Venezuela Snouted Treefrog

small humid plastic bags, transported from the collection site to the field laboratory with a walk of no longer than one hour. They were kept in captivity until the next morning, when the fixation procedures were carried out. Individuals were euthanized with 0.5% xylocaine according to regulations of the Federal Council of Veterinary Medicine (Law 5.517/68 article 16, "f"). The animals were fixed in 10% formaldehyde and subsequently maintained in 70% (v/v) alcohol. At the end of field expedition, the treefrogs were taken to the laboratory, to undergo histological analysis.

For light microscopic (LM) analysis, 3–5 mm thick sections from the ventral, ventrolateral and dorsal regions of the integument were processed according to standard histological techniques for paraffin embedding before sectioning, i.e., the sections were quickly washed in water, dehydrated (70%, 90%, twice in 100% ethanol; 30 minutes each), clarified twice in xylene (30 minutes each), infiltrated and embedded in paraffin. The 5-µm thick serial histological sections were stained with hematoxylin-eosin (HE) (Lillie and Fulmer 1976), which is the standard stain for histological examination of animal tissues, staining the nuclei and cytoplasm in blue with the extracellular matrix in pale pink. Mallory's trichrome (Lillie and Fulmer 1976) was used since it is a good stain for distinguishing cellular from extracellular elements and is especially suitable for studying connective tissue, staining the collagenous fibers in blue, red blood cells in orange, and nuclei in red. Staining with 1% Alcian blue (AB) 8GX at pH 2.5 (Mowry 1963, Kiernan 1990) was utilized to detect sulfated and carboxylated glycoconjugates (in light blue). In addition, HE-stained slices were observed under a light microscope using polarized light in order to detect iridophores (an iridescent cell), since these cell types possess reflective structures (Bagnara et al. 1968) that can be easily recognized under polarized light. Sections were analyzed using a Leica DM750 microscope, and the images were captured using a Leica DFC452 digital camera.

RESULTS

In general, the integuments of all species showed the basic morphological structure of the anuran integument, essentially



Figures 1–10. Photograph of the species: (1) *O. albicans*; (2) *O. angrensis*; (3) *O. flavoguttata*; (4) *S. hayii*; (5) *O. humilis*; (6) *O. perpusilla*; (7) *S. similis*; (8) *O. trapicheroi*; (9) *O. v-signata*; (10) *S. x-signatus*.

being formed by an epidermal and dermal layer (Figs 11–55).

The epidermis was relatively thin and consisted of a partially keratinized stratified squamous epithelium supported by a dermis. Although the keratinocyte was the predominant cell type, native flask cells were also visualized. In addition, epidermal cells were organized into a basal layer, an intermediate layer and an outermost layer. The epidermal cells of the outermost layer were partially keratinized, since their nuclear profiles could be easily visualized.

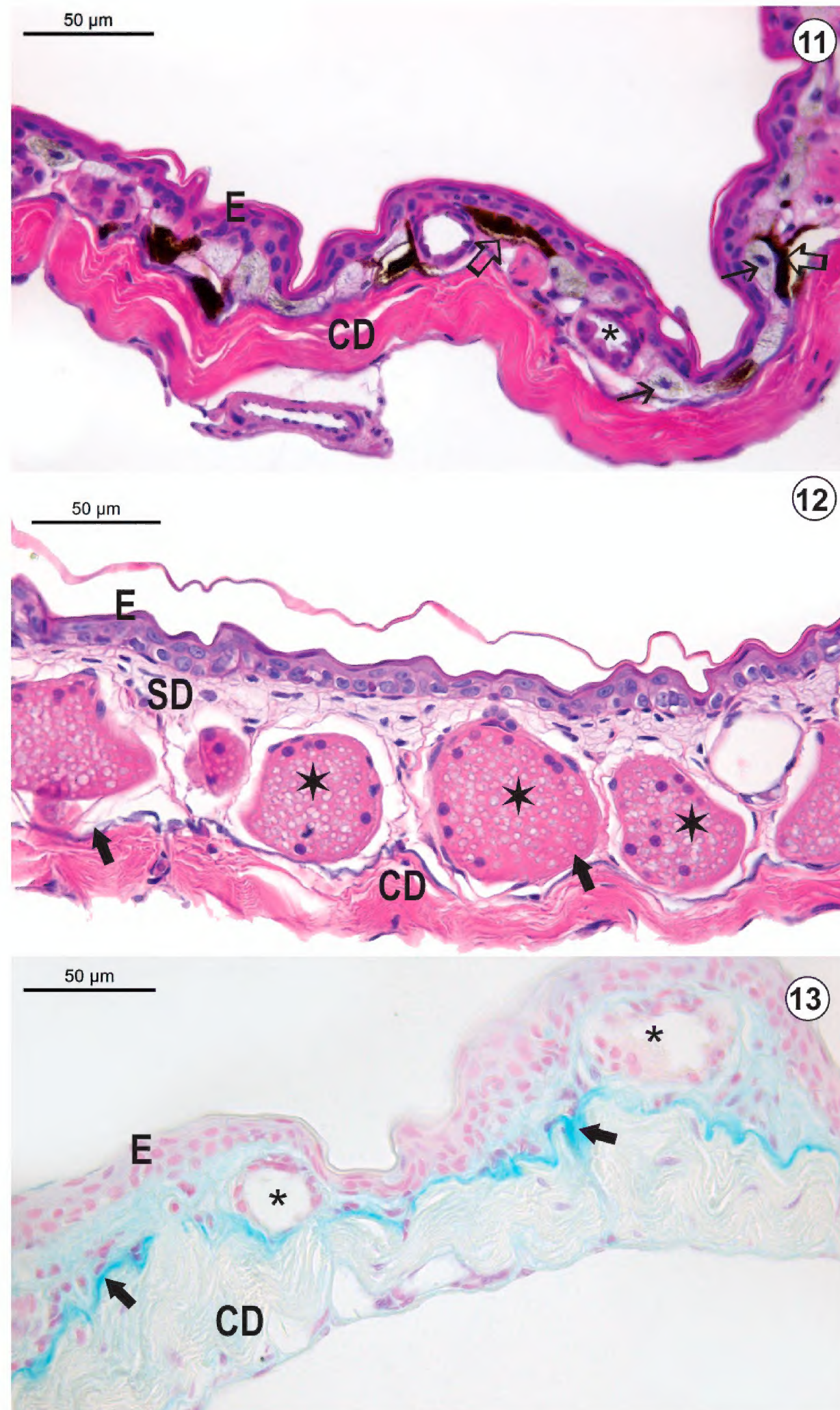
Considering the number of epidermal cell layers of the integument, subtle differences between species and body regions were observed (Table 2), i.e., the thicker epidermis was observed in *Ololygon humilis* (Lutz & Lutz, 1954) and *Scinax x-signatus* (Spix, 1824).

The dermis was subdivided into the spongy dermis, composed of loose connective tissue, and the compact dermis, which rested on the hypodermis. Moreover, the compact dermis was formed by collagenous fibers organized in a series of alternating layers, compactly arranged in a crisscross manner. Between the spongy dermis and the compact dermis, irregular basophilic deposits occurred scattered through this boundary region, corresponding to the Eberth-Katschenko (EK) layer.

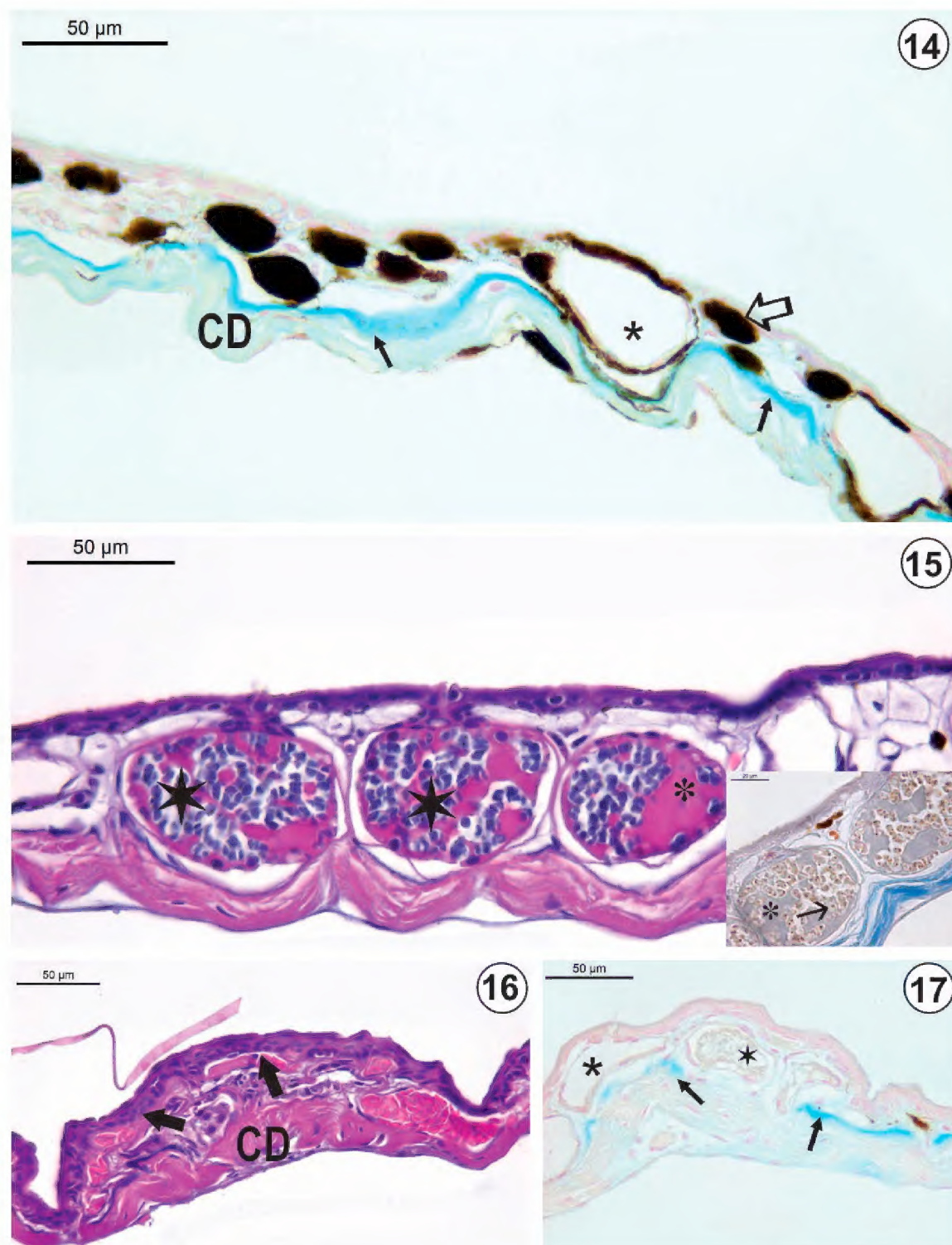
Table 2. Number of epithelial cell layers of *Ololygon* spp. and *Scinax* spp.

	Number of epithelial cell layers	
	Dorsal	Ventral
<i>O. albicans</i>	4	4–5
<i>O. angrensis</i>	4	4–5
<i>O. flavoguttata</i>	4	4–5
<i>O. humilis</i>	4	8
<i>O. perpusilla</i>	3–4	4–5
<i>O. trapicheroi</i>	5	5
<i>O. v-signata</i>	4	4
<i>S. similis</i>	4	4
<i>S. hayii</i>	4	4–5
<i>S. x-signatus</i>	5	7

Another typical feature of these treefrogs was the occurrence of different exocrine glands, whose secretory portions were housed in the spongy dermis (Figs 11–55). From the secretory portion, a single unbranched excretory duct passed over the epidermis and opened on the integument surface. These ducts were lined by two layers of cuboid cells.



Figures 11–13. Light micrograph of the integument of *O. albicans*: (11) Dorsal region (HE-staining); (12) Ventrolateral region (HE-staining); (13) Ventral region (AB-method). In all integument regions, the epidermis (E) rests on the dermis, which is subdivided into the spongy dermis (SD) and the compact dermis (CD). Iridophores (→) occur in the dorsal region; however, they are absent in both ventrolateral and ventral regions. Melanophores (⇒) in the spongy dermis. Both serous (*) and apocrine glands (*) occur in the spongy dermis. No glandular cell reacts to AB-method, suggesting that secretory units is made up of serous cells. The EK-layer (➔) exhibits its typical basophilic staining. Note clusters of apocrine glands (*) with heterogeneous content in the ventrolateral integument. The EK-layer (➔) exhibits typical alcianophilic reaction of its glycoconjugate content. Large blood vessels occur in the hypodermis.

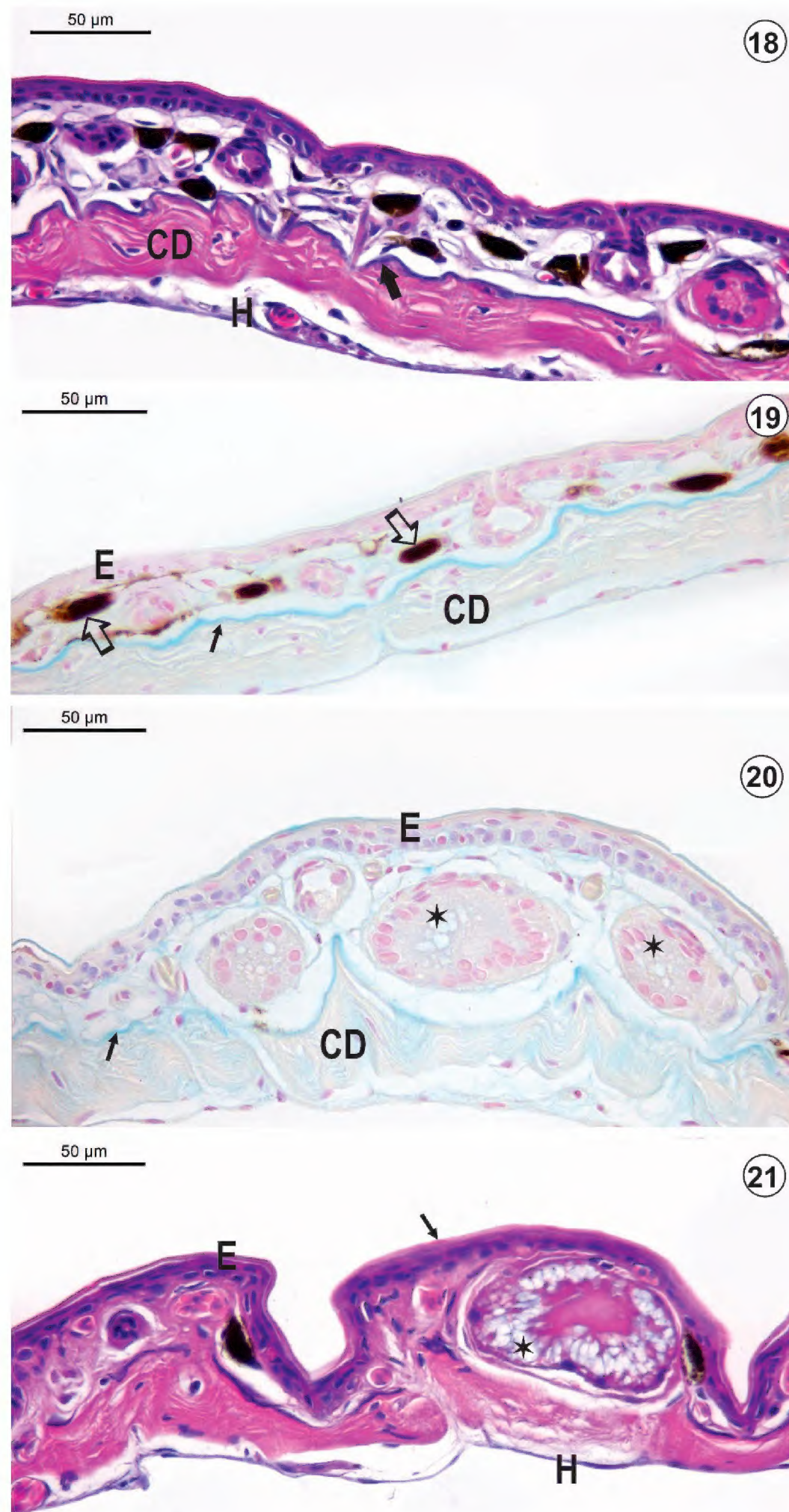


Figures 14–17. Light micrograph of the integument of *O. angrensis*: (14) Dorsal region (AB-staining); (15) Ventrolateral region (HE-staining), inset (Mallory's trichrome staining); (16) Ventral region (HE-staining); (17) Ventral region (AB-method). The melanophores (⇔) are numerous and located just beneath of the epidermis as well as around de glandular secretory units. They occur also in the hypodermis, but absent in the ventral integument. Iridophores (→) occur in the spongy dermis of the dorsal region just beneath the epidermis. No iridophore is visualized in the ventral region of the integument. Clusters of apocrine glands (*) with heterogeneous intake predominate at ventrolateral integument; inset of Fig. 15: Observe the granular content with dense stained core (➔) intermingled with cytoplasmic material (*). The EK-layer (➔) is continuous in the ventral region, but discontinuous in the ventral integument. Serous glands (*) occur in all body regions. Note blood vessels (➔) in the spongy dermis. CD = compact dermis.

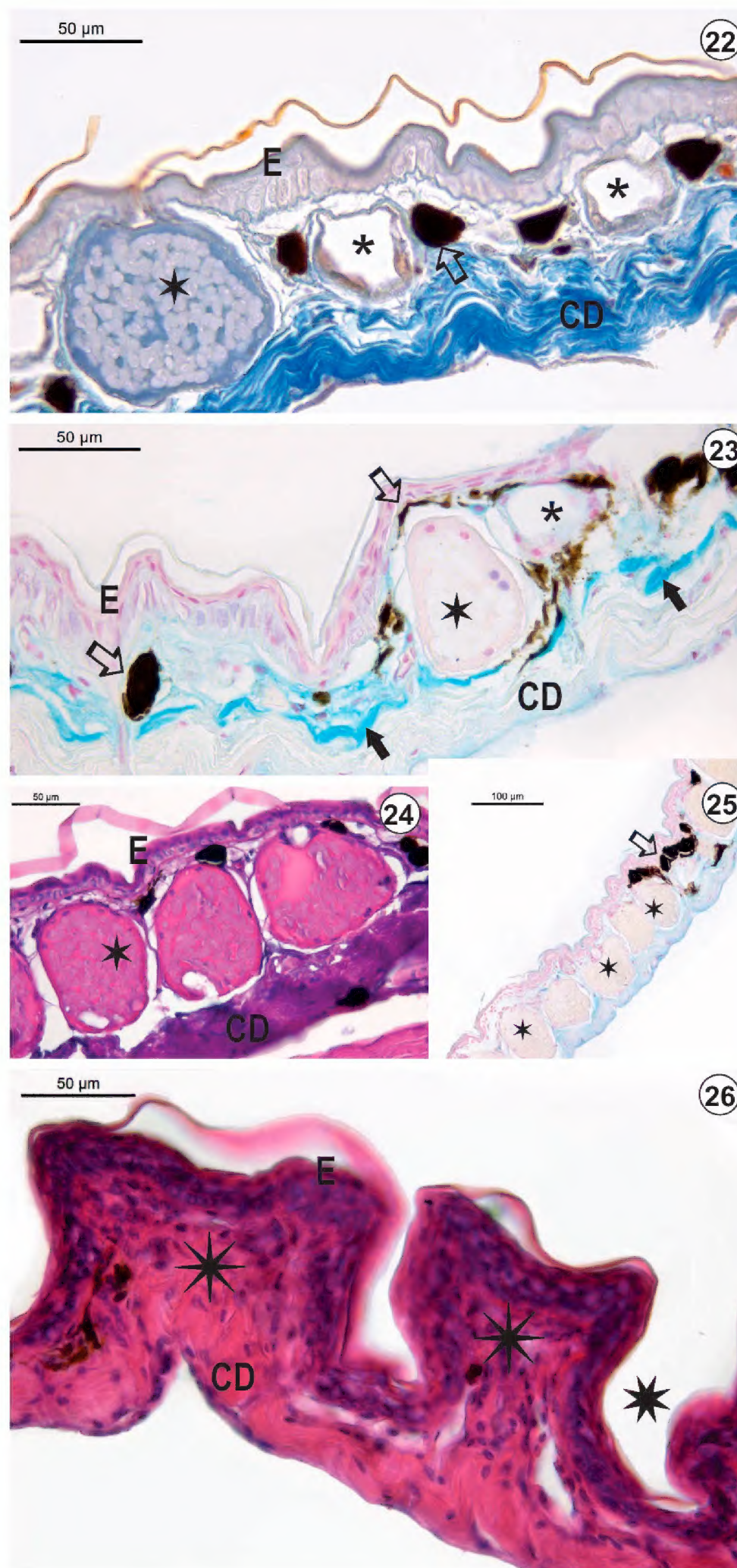
Depending on the dye affinity to their secretion, the merocrine glands of the serous and mixed types were visualized. Serous glands are formed by only serous-secreting cells, which have spherical nuclei and an acidophilic cytoplasm. Mixed secretory units are made up of mucous cells (nuclei generally flattened and displaced to the basal portion of the secretory cells and basophilic cytoplasm) and serous cells. It is noteworthy that mucous glands essentially formed by mucous cells did not occur in the integuments of all tree frogs (Table 3 and Figs 11–55).

While serous glands were visualized in all species, mixed glands occurred in *Ololygon albicans* (Bokermann, 1967), *O. angrensis*, *O. flavoguttata* (Lutz & Lutz, 1939), *O. humilis*, *O. trapicheroi* (Lutz & Lutz, 1954), *O. v-signata* (Lutz, 1968), *Scinax similis* (Cochran, 1952) and *S. x-signatus*, but they did not occur in *S. hayii* (Barbour, 1909) and *O. perpusilla* (Lutz & Lutz, 1939) (Table 3).

Considering the apocrine glands, their secretory portion consisted of syncytial units, varying according to their content and dye affinity. Their secretory products were made up of small



Figures 18–21. Light micrograph of the integument of *O. flavoguttata*: (18) Dorsal region (HE-staining); (19) Dorsal region (AB-method); (20) Ventrolateral region (HE-staining) (21) Ventral region (HE-staining). The epidermis (E) is partially keratinized, and the outermost cell layer exhibits the nuclear profiles (→). Melanophores (⇨) occur in the spongy dermis of both dorsal and ventrolateral regions; they are more frequent in the dorsal integument. Small clusters of apocrine glands with heterogeneous content occur in the ventrolateral integument. In the ventral region, the lipid content (★), mixed with basophilic material, is easily visualized in the apocrine glands (Fig. 21). In the dorsal region, the EK-layer (➔) is continuous, but discontinuous in the ventrolateral integument, being less alcianophilic in the ventral region when compared to other integument regions. CD = compact dermis; H = hypodermis.



Figures 22–26. Light micrograph of the integument of *S. hayii*: (22) Dorsal region (Mallory's trichrome staining); (23) Dorsal region (AB-method); (24) Ventrolateral region (HE-staining) (25) Ventral region (AB-method); (26) Ventral region (HE-staining). In the dorsal integument, exocrine glands are more frequent, mainly the serous glands (★). Melanophores (⇒) occur in the spongy dermis, even around the secretory portion of glands. Note clusters of apocrine glands with heterogeneous content (★) in the spongy dermis of the ventrolateral integument. The EK-layer (⇒) is continuous in the dorsal integument, but absent in the ventral region. Slight cutaneous elevations (★) in the ventral integument are formed by the epidermis and the dermis, mainly the spongy dermis. They are separated by grooves (★). CD = compact dermis.

Table 3. Occurrence of exocrine merocrine glands in the integument of *Oloygon* spp. and *Scinax* spp.

	Merocrine glands								
	Mucous gland			Serous gland			Mixed glands		
	Dorsal	Ventrolateral	Ventral	Dorsal	Ventrolateral	Ventral	Dorsal	Ventrolateral	Ventral
<i>O. albicans</i>	–	–	–	+	+	+	–	–	+
<i>O. angrensis</i>	–	–	–	+	+	+	–	+	–
<i>O. flavoguttata</i>	–	–	–	+	+	+	–	+	–
<i>O. humilis</i>	–	–	–	+	+	+	+	+	+
<i>O. perpusilla</i>	–	–	–	+	+	–	–	–	–
<i>O. trapicheroi</i>	–	–	–	+	+	+	+	+	+
<i>O. v-signata</i>	–	–	–	+	–	–	+	+	+
<i>S. similis</i>	–	–	–	+	+	+	+	+	+
<i>S. hayii</i>	–	–	–	+	+	+	–	–	–
<i>S. x-signatus</i>	–	–	–	+	–	–	+	+	+

(+) Present, (–) absent.

Table 4. Occurrence of apocrine glands in the integument of *Oloygon* spp. and *Scinax* spp.

	Heterogeneous content					
	Basophilic			Lipid		
	Dorsal	Ventrolateral	Ventral	Dorsal	Ventrolateral	Ventral
<i>O. albicans</i>	+	+/x	+	+	+	+
<i>O. angrensis</i>	+	–	–	+	+	–
<i>O. flavoguttata</i>	+	+/x	+	–	+	+
<i>O. humilis</i>	+	+	+	+	+	+
<i>O. perpusilla</i>	+	+	–	+	+	–
<i>O. trapicheroi</i>	+	+	+	+	–	–
<i>O. v-signata</i>	+	+	+	+	+	+
<i>S. similis</i>	–	–	–	+	+	–
<i>S. hayii</i>	+	+/x	–	+	+	+
<i>S. x-signatus</i>	–	+	–	+	+	–

(+) Present, (–) absent, (x) occurrence in clusters.

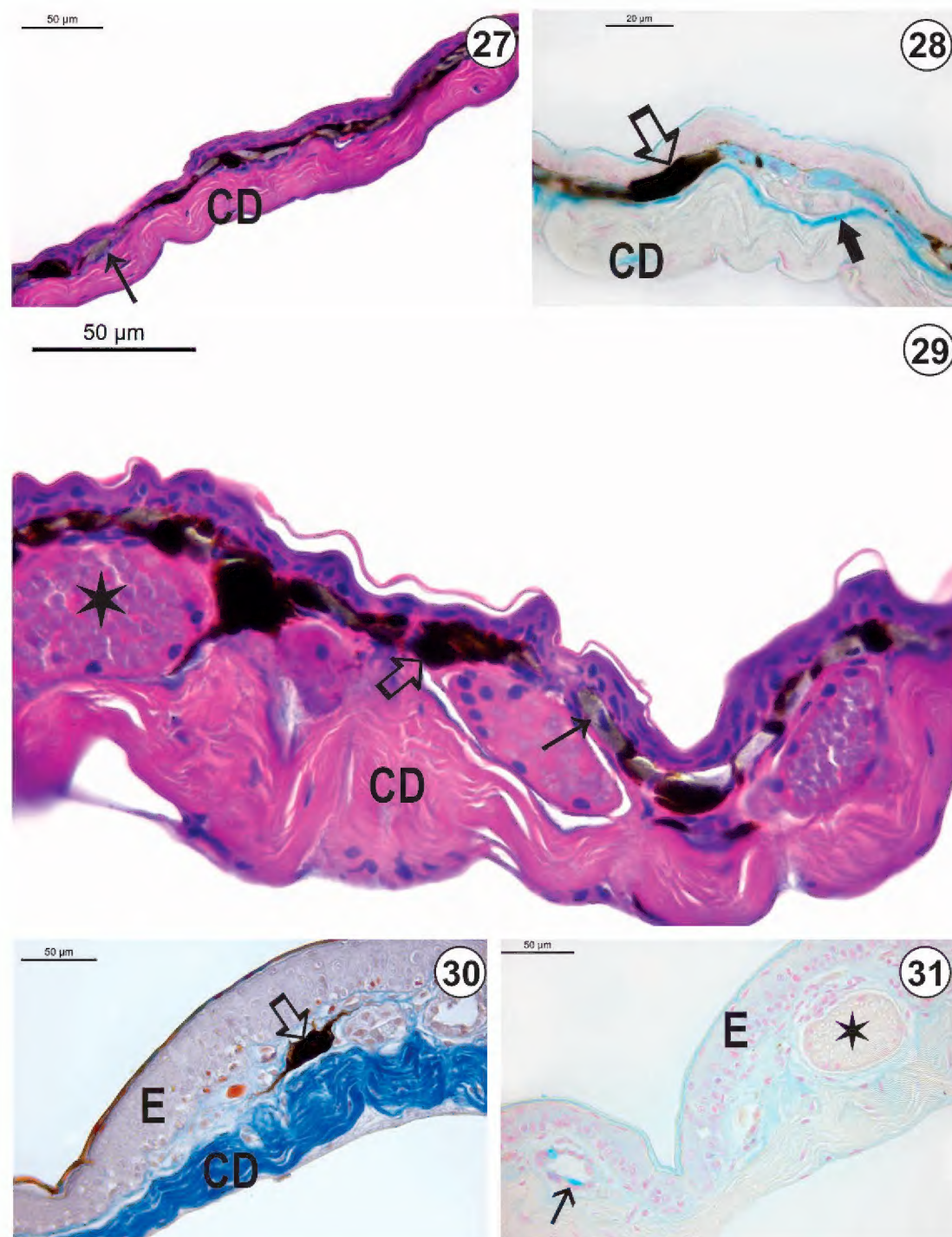
acidophilic granules, seemingly due to their protein content. Nevertheless, some apocrine glands revealed that this secretion is a mixture of basophilic and lipid contents, intermingled with cytoplasmic material; their rounded nuclei were displaced to the cell basal domain. In some glands, the secretory product was constituted of acidophilic cytoplasmic material mixed with heterogeneous material, which revealed slight basophilic reaction usually associated with lipid material (Table 4).

In *O. angrensis*, the secretory product of the apocrine glands exhibited basophilic granules with an acidophil core after staining with Mallory’s trichrome (Figs 14–17). Furthermore, these secretory granules were also observed in *S. similis* (Figs 36–40) and contained lipids associated with basophilic material. We suggest that the basophilic staining was due to carbohydrates of glycolipids, since they showed no alcianophilic reaction typical of acid polysaccharides.

Table 5. Occurrence of pigment cells in the integument of *Oloygon* spp. and *Scinax* spp.

	Melanophores		Iridophores	
	Dorsal	Ventral	Dorsal	Ventral
<i>O. albicans</i>	–/+	–/+	+	+
<i>O. angrensis</i>	+	–/+	–/+	–
<i>O. flavoguttata</i>	+	–/+	–	–
<i>O. humilis</i>	+	–/+	+	–
<i>O. perpusilla</i>	+	–	–	–
<i>O. trapicheroi</i>	–/+	–	+	–
<i>O. v-signata</i>	+	–	–	–
<i>S. similis</i>	+	–/+	+	+
<i>S. hayii</i>	+	–/+	–	–
<i>S. x-signatus</i>	+	–	–	–

(+) Widespread, (–/+) occasional, (–) not visualized.



Figures 27–31. Light micrograph of the integument of *O. humilis*: (27) Dorsal region (HE-staining); (28) Dorsal region (AB-method); (29) Ventrolateral region (HE-staining) (30) Ventral region (Mallory's trichrome staining); (31) Ventral region (AB-method). In the dorsal region, the spongy dermis is poorly developed. Melanophores (⇨) are visualized in all integument regions; however, iridophores (→) are visualized only in both dorsal and ventrolateral integument. Both pigment cells are located just beneath the epidermis. Alcianophilic reaction is observed in cytoplasm of iridophores as well as in the EK-layer (⇨) of the dorsal integument. The EK-layer is absent in the ventral integument. Apocrine glands with heterogeneous content (★) occur in both ventrolateral and ventral integument. In *S. humilis*, mixed glands (→) are visualized in the ventral region, being formed by serous and mucous cells. Mucous cells exhibit alcianophilic reaction. E = epidermis; CD = compact dermis.

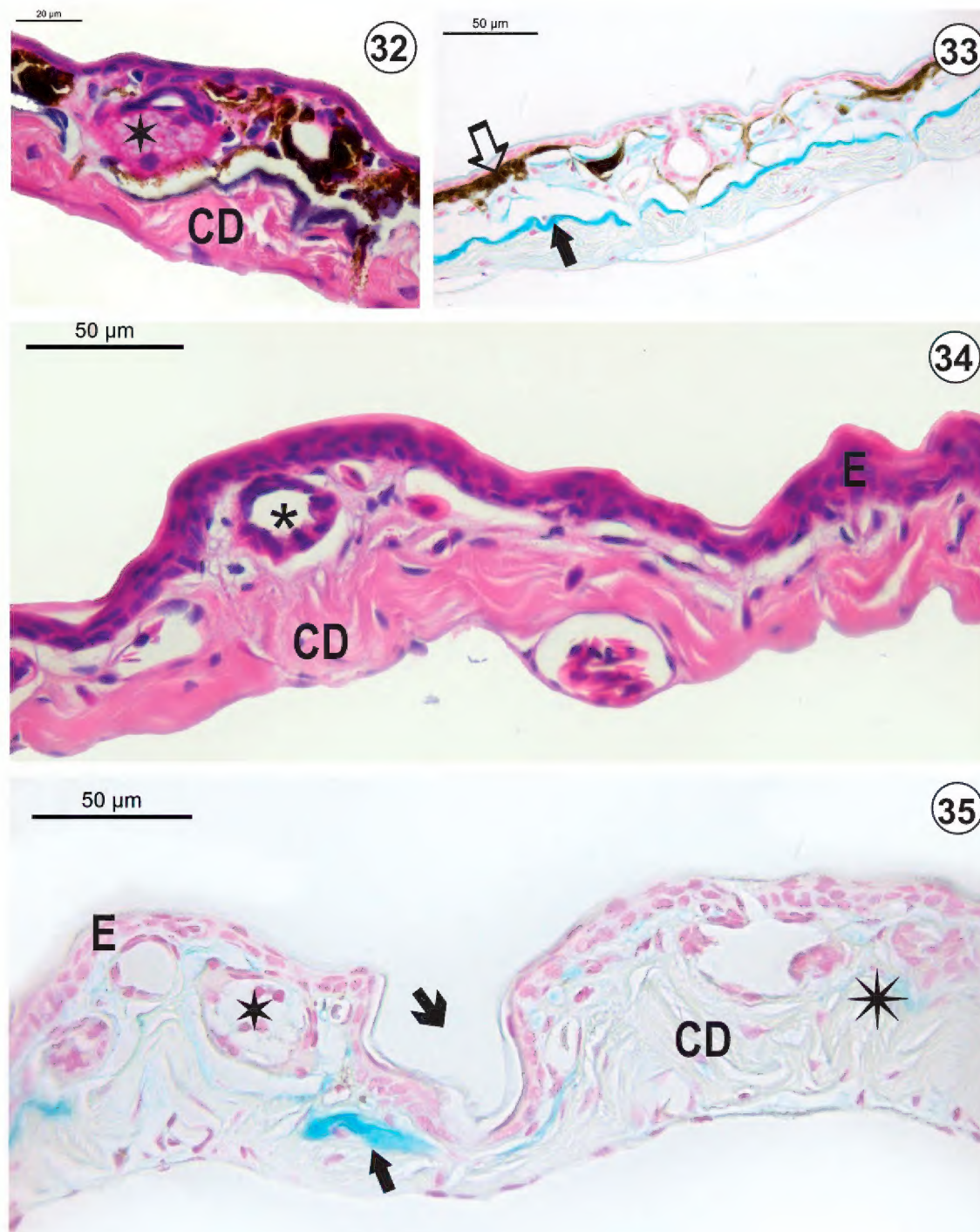
In *O. albicans*, *O. angrensis*, *O. flavoguttata* and *S. hayii*, apocrine glands with heterogeneous content were more frequent in the ventrolateral integument, occurring as small clusters (Figs 11–26).

Pigment cells, such as melanophores and iridophores, were identified in the integument, occurring in the spongy dermis, just beneath the basal lamina. While melanophores were identified through the typical brownish color of their melanin granules under light microscopy, iridophores were visualized by polarized light microscopy through their reflective or iridescent pigments (Figs 11–55). Melanophores also occurred in the hypodermis of *O. angrensis*, *O. v-signata* and *S. similis* (Figs 4–11, 36–40, 52–55),

while iridophores were only detected in the hypodermis of *S. similis*. The occurrence of both pigment cells varied according to the species and integument region (Table 5).

The Eberth-Katschenko (EK) layer occurred between the spongy and compact dermis and was recognized through its typical basophilic and alcianophilic reaction after using the HE- and AB-methods, respectively (Table 6; Figs 11–55). The EK-layer was visualized in the dorsal region of the integument of all *Oloygon* species, but absent in all species of *Scinax*.

Cutaneous elevations occurred in the ventral integument of *O. angrensis*, *O. flavoguttata*, *O. perpusilla*, *S. similis*, *O.*



Figures 32–35. Light micrograph of the integument of *O. perpusilla*: (32) Dorsal region (HE-staining); (33) Dorsal region (AB-method); (34) Ventrolateral region (HE-staining) (35) Ventral region (AB-method). The spongy dermis houses both apocrine glands with heterogeneous content (*) and serous glands (*). Melanophores (⇨) occur in the dorsal integument, but they are not identified in either the ventrolateral or ventral region. Iridophores did not occur in all integument regions. The EK-layer (➔) is a well defined continuous layer occurs as irregular deposits between the spongy and compact dermis of the ventral region. Cutaneous elevations (*) are separated by grooves (➔) in the ventral region. CD = compact dermis.

trapicheroi and *S. x-signatus* (Figs 14–21, 32–40, 52–55). These slight elevations were formed by the epidermis, followed by the dermis, being separated by grooves.

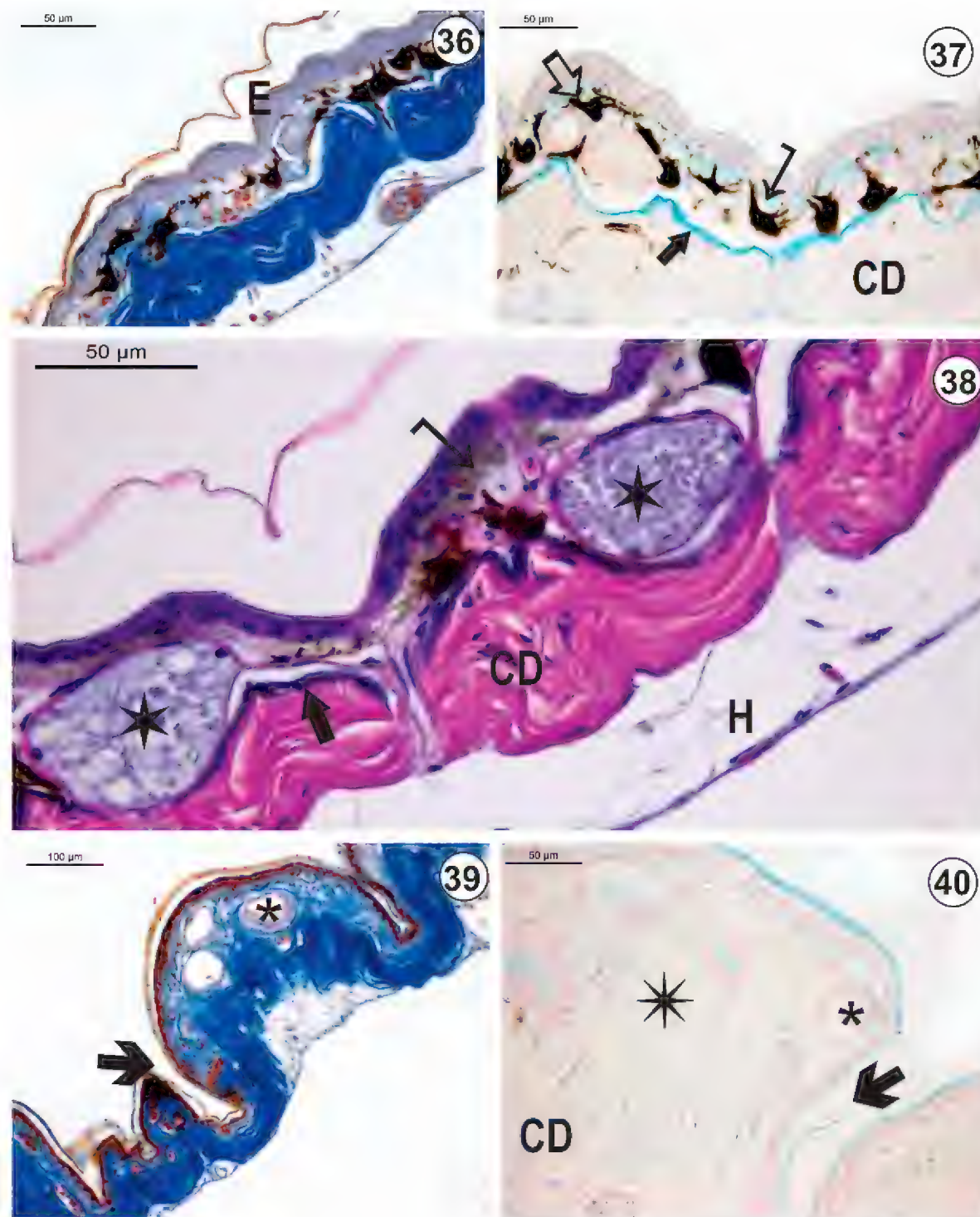
DISCUSSION

Although some studies of the anuran integument are available, such as those of bufonids (de Brito-Gitirana and Azevedo 2005, de Brito-Gitirana et al. 2007, Almeida et al. 2007, Felsemburgh et al. 2009), ranids (Azevedo et al. 2006, Pelli et al. 2010), and leptodactylids (Goniakowska-Witalinska and Kubiczek 1998, Warburg et al. 2000, Nosiet al. 2002, Barbeau and Lillywhite 2005, Felsemburgh et al. 2007, Gonçalves and de

Brito-Gitirana 2008, Rigolo et al. 2008), information about the integument structure of hylids is still insufficient.

In this study, the integument showed the basic structure as already described for other anurans, i.e., the epidermis rests on a dermis, which is divided into a spongy and a compact dermis. The majority of anurans display this structural pattern of the integument (Elkan 1968, Goniakowska-Witalinska and Kubiczek 1998, Warburg et al. 2000, de Brito-Gitirana and Azevedo 2005, Delfino et al. 2006, Felsemburgh et al. 2007, Gonçalves and de Brito-Gitirana 2008, Felsemburgh et al. 2009).

In general, the epidermis varied from 4–5 cell layers. However, in *O. humilis* and *S. x-signatus* the ventral epidermis was thicker, and it may be related to species habitat. Nevertheless,

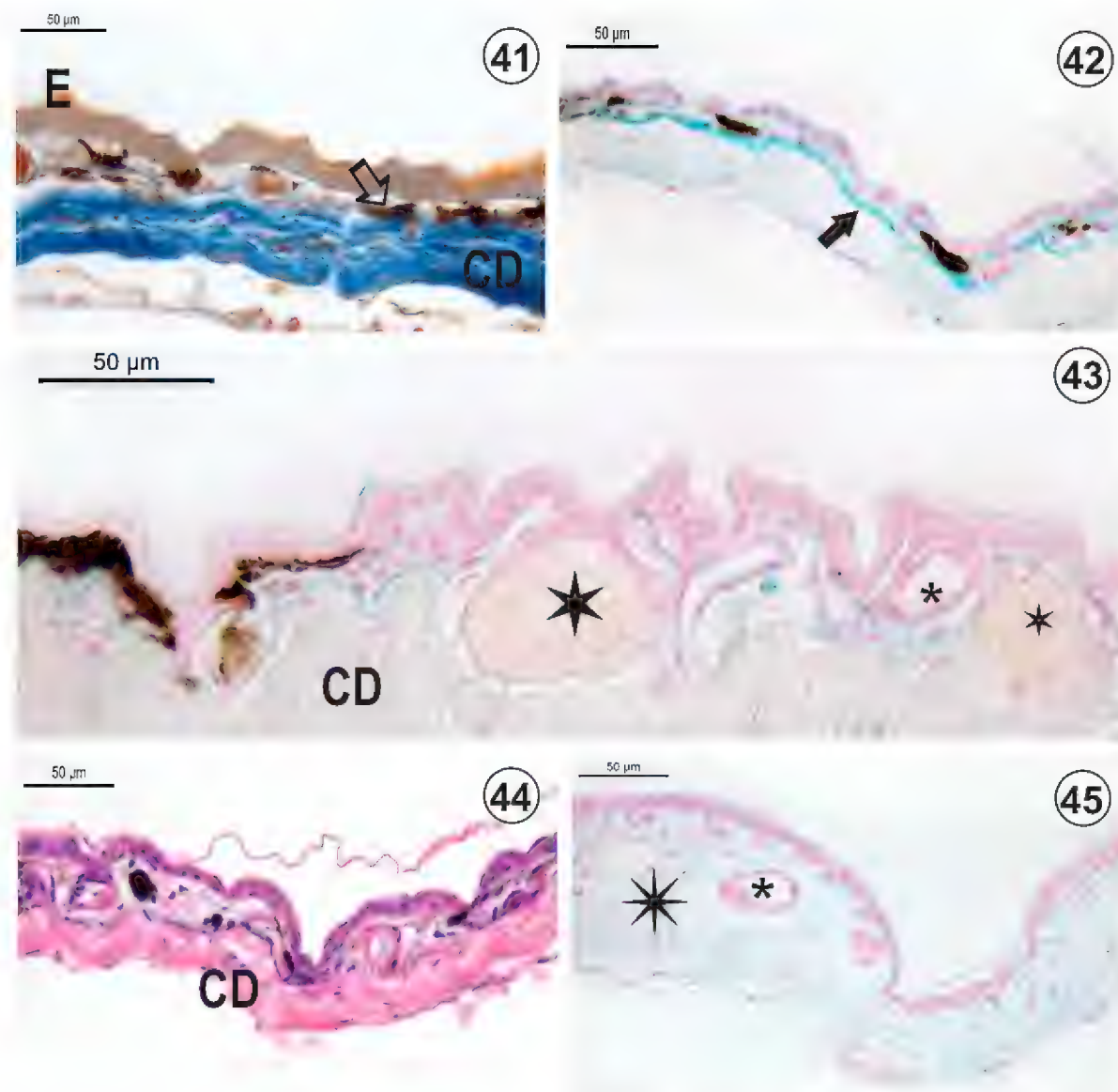


Figures 36–40. Light micrograph of the integument of *S. similis*: (36) Dorsal region (Mallory's trichrome staining); (37) Dorsal region (AB-method); (38) Ventrolateral region (HE-staining); (39) Ventral region (Mallory's trichrome staining); (40) Ventral region (AB-method). Melanophores (⇒) and iridophores (→) organized as chromatophore units occur in both dorsal and ventrolateral integument. Iridophores exhibit alcianophilic reaction. Note serous glands (★) and apocrine glands with granular content (★) in the spongy dermis. In the ventral region, cutaneous elevations (★) are separated by prominent grooves (→). The EK-layer (⇒) occur in the dorsal integument but not in the ventral integument. Moreover, the epidermis (E) of the ventral region is more developed than other integument regions. E = epidermis; CD = compact dermis; H = hypodermis.

we did not find detailed behavioral data in the literature to support this explanation. In mammals, in some body areas, friction and other forces dictate the thickness of the lining epithelium, since the number of epithelial cell layers is related to epithelial resistance (Ham 1977, de Brito-Gitirana 2015).

Anuran glands have received significant attention. They have been described as being of different types, like mucous, serous, lipid (or wax), and mixed (seromucous) glands. However, some authors have named the granular glands as poison or serous glands (Mills and Prum 1984, Duellmann and Trueb 1994, Brizzi et al. 2002).

In this study, well-established histological criteria to categorize the cutaneous gland of mammals (Ham 1977, Kierzenbaum 2004) were used in order to adopt a coherent histological classification, especially in reference to anuran glands, since they exhibit variable morphology. Thus, on the basis of how their secretory products are released to the external environment, the exocrine glands can be classified as holocrine, apocrine or merocrine. Holocrine glands release both secretions and entire cells, while apocrine glands release the secretory product and cytoplasmic matrix of the apical portion of the cell. In merocrine cells, no cytoplasm is lost and the gland uses exocytosis



Figures 41–45. Light micrograph of the integument of *O. trapicheroi*: (41) Dorsal region (Mallory's trichrome staining); (42) Dorsal region (AB-method); (43) Ventrolateral region (AB-method); (44) Ventral region (HE-staining); (45) Ventral region (AB-method). Melanophores (⇒) occur in the spongy dermis of the dorsal integument, and in the ventrolateral region as isolated groups. They are absent in the ventral integument. In the dorsal region, the EK-layer (⇒) is continuous and well stained by the AB-method. Isolated serous glands (★) occur in all integument regions. The apocrine glands with granular content (✱) are visualized in the ventrolateral integument, where they are more developed. In the ventral region, cutaneous elevations (✱) are also separated by grooves. The dermis contains several small blood vessels. E= epidermis; CD = compact dermis.

Table 6. Occurrence of the EK-layer in the integument of *Oloygon* spp. and *Scinax* spp. The EK-layer can occur as a continuous or discontinuous layer.

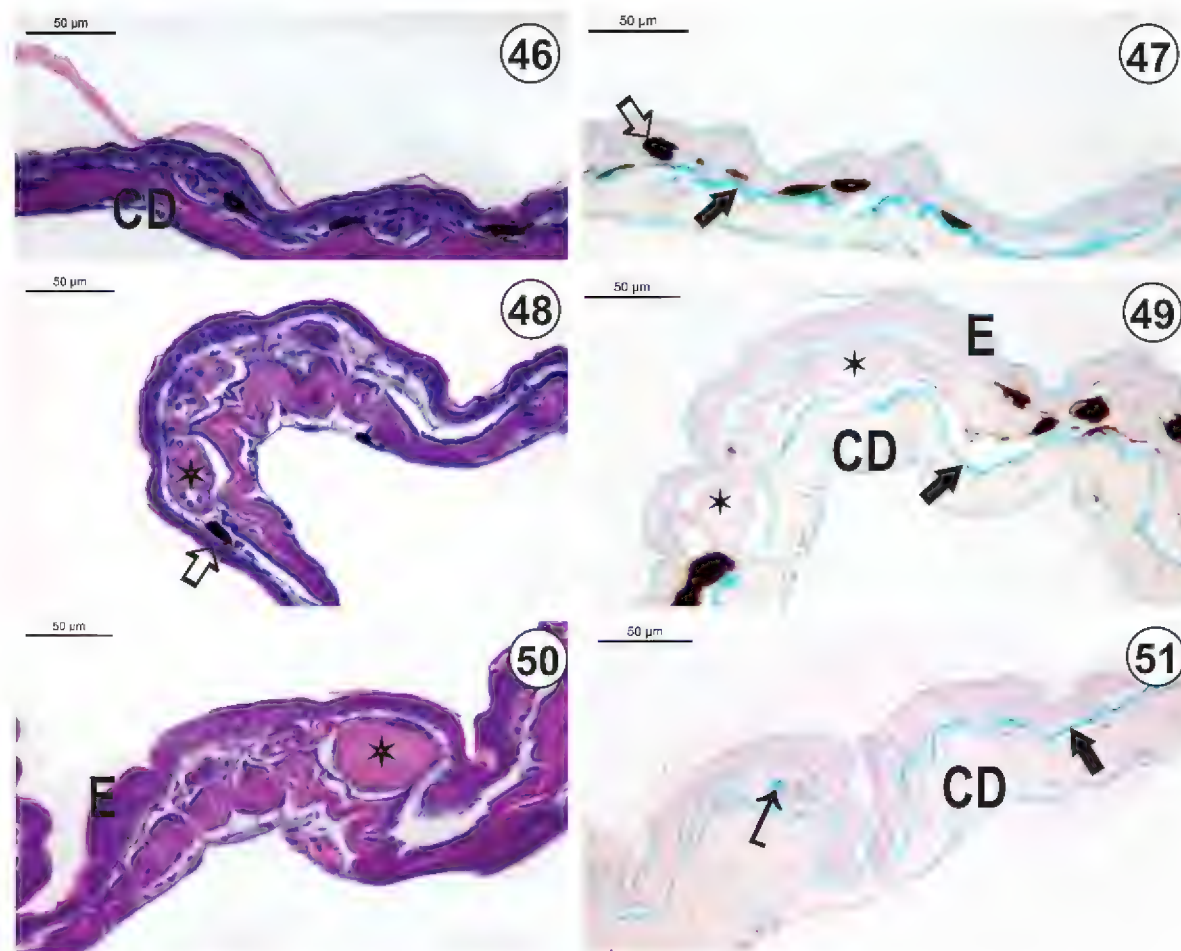
	EK-layer		
	Dorsal	Ventrolateral	Ventral
<i>O. albicans</i>	+/Ø	+/Ø	+/Ø
<i>O. angrensis</i>	+/Ø	+/Ø	+/â
<i>O. flavoguttata</i>	+/Ø	+/â	+/Ø
<i>O. humilis</i>	+/Ø	+/â	–
<i>O. perpusilla</i>	+/Ø	–	+/â
<i>O. trapicheroi</i>	+/Ø	–	–
<i>O. v-signata</i>	+/Ø	+/Ø	+/Ø
<i>S. similis</i>	+/Ø	+/Ø	–
<i>S. hayii</i>	+/â	–	–
<i>S. x-signatus</i>	+/Ø	+/Ø	–

(+) Present, (–) absent, (Ø) continuous, (â) discontinuous.

to release its secretory product to the extracellular space (Ham 1977, Kierzenbaum 2004).

Given that the secretory cell remains intact, according to the type of secretion produced, the merocrine gland of mammals is classified as serous, mucous or mixed. Serous glands are essentially composed of serous cells with large spherical nuclei and an acidophilic cytoplasm that is individualized by the cell membrane. Mucous glands are composed by the mucus-secreting cells, which exhibit an irregular shape; the nuclei are basally located, and their cytoplasm is basophilic (through HE-staining), alcianophilic (through AB-staining to detect acid glycoconjugates), and/or PAS positive (detect neutral glycoprotein). Mixed glands are made up of both serous and mucous cells constituting the same secretory portion (Ham 1977, Kierszenbaum 2004, de Brito-Gitirana 2013).

In some anurans, serous and granular glands have been considered as the same type. Nevertheless, various subtypes of



Figures 46–51. Light micrograph of the integument of *O. v-signata*: (46) Dorsal region (HE-staining); (47) Dorsal region (AB-method); (48) Ventrolateral region (HE-staining); (49) Ventrolateral region (AB-method); (50) Ventral region (HE-staining); (51) Ventral region (AB-method). Melanophores (⇒) occur in the spongy dermis that is poorly developed in the dorsal region. The EK-layer (⇒) is a continuous layer in all integument regions. Apocrine glands (*) with heterogeneous content occur in both ventrolateral and ventral regions. Serous glands are visualized in both dorsal and ventrolateral integument. Mixed glands (•) are observed in the ventral region. E = epidermis; CD = compact dermis.

serous glands with high morphological variability have been reported (Goniakowska-Witalinska and Kubiczek 1998, Delfino et al. 1999, Warburg et al. 2000, Nosi et al. 2002, Brunetti et al. 2012, Moreno-Gómez et al. 2014).

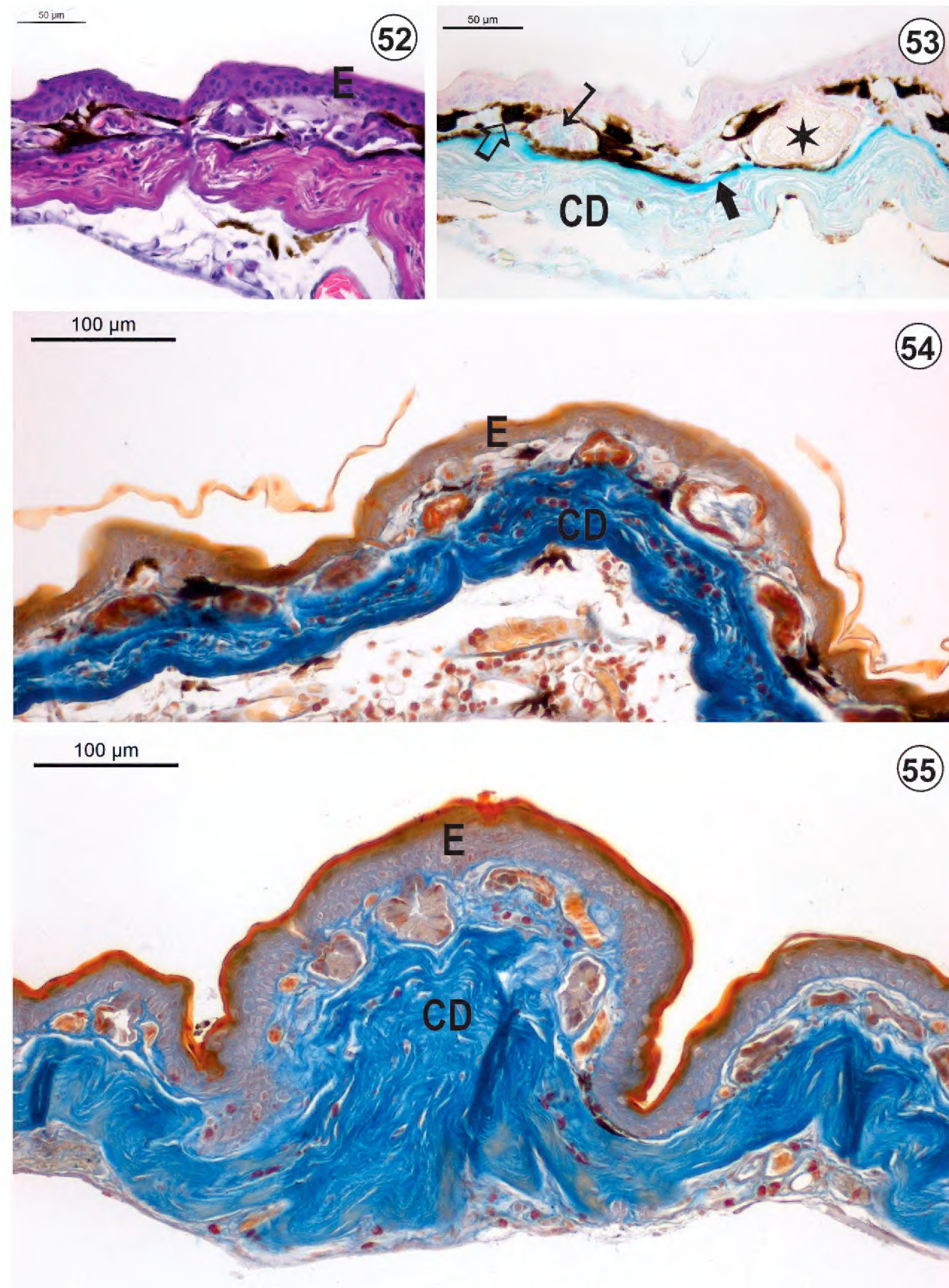
Actually, the granular gland described by Fox (1986a, b) and Duellman and Trueb (1994) is a kind of apocrine gland since its secretory product is a mixture of cytoplasm and secretory product. Furthermore, this glandular type is composed of a syncytium since the large mass of cytoplasm with many nuclei is not separated into individual cells by a cell membrane. In fact, the apocrine glands of anurans exhibit a high level of polymorphism.

In bufonids (de Brito-Gitirana and Azevedo 2005, de Brito-Gitirana et al. 2007, Almeida et al. 2007, Felsemburgh et al. 2009), ranids (Azevedo et al. 2006, Pelli et al. 2010), and leptodactylids (Felsemburgh et al. 2007, Gonçalves and de Brito-Gitirana 2008), apocrine granular glands occur scattered throughout the dorsal and ventral integument, exhibiting acidophilic granular secretion. In the integument of *S. similis*, the apocrine granular gland resembles those Ib type glands described for *Engystomops pustulosus* (Cope, 1864) (Delfino et al. 2015). In addition, apocrine glands with heterogeneous content occurred

in the integument of all species of Scinaxinae. Nevertheless, in *O. angrensis*, the secretory product contained peculiar basophilic granules with dense cores intermingled with acidophilic cytoplasm, and their histological features resembled those of type Ia glands described for *Pithecopus azureus* (Cope, 1862) (former *Phyllomedusa azurea*) (Nosi et al. 2002).

In all species, the apocrine gland included an acidophilic content mixed with slightly basophilic material. Although the secretory granules exhibited basophilic affinity, they revealed no alcianophilic reaction, demonstrating that their content had no glycoconjugate, and this material probably consists of glycolipids. On the other hand, lipids have been observed in cutaneous secretions of phyllomedusine (Blaylock et al. 1976), hylids *Litoria fallax* (Peter, 1880) and *L. peronii* (Tschidi, 1838) (Amey and Grigg 1995), African frogs (Withers et al. 1984), *Ranoidea australis* (Gray, 1842) (former *Cyclorana australis*) (Christian and Parry 1997), and the hylid *Ranoidea caerulea* (White, 1790) (former *Litoria caerulea*) (Warburg et al. 2000).

According to Warburg and co-workers (2000), lipid content represents the main adaptation of xeric-inhabiting arboreal frogs, enabling them to remain exposed throughout the year, even during dry seasons. Cutaneous lipids in tree frogs – *Phyl-*



Figures 52–55. Light micrograph of the integument of *S. x-signatus*: (52) Dorsal region (HE-staining); (53) Dorsal region (AB-method); (54) Ventrolateral region (Mallory's trichrome staining); (55) Ventral region (Mallory's trichrome staining). The epidermis (E) is slightly thicker when compared to those of other hylids, as in the compact dermis (CD). Melanophores (⇒) are visualized in both dorsal and ventrolateral integument just beneath the epidermis. Serous glands are present in all regions; however, some of them show slightly alcianophilic content (→) in both ventrolateral and ventral regions. The apocrine glands (★) with granular content occur in both dorsal and ventrolateral regions. The EK-layer (➡) is visualized in both dorsal and ventrolateral integument but is absent in the ventral integument.

lomedusa sauvagei Boulenger, 1882, *P. iherengii* Boulenger, 1885, *P. boliviana* Boulenger, 1902, *Ranoid gracilentia* (Peters, 1869), *R. caerulea* (White, 1790), *Polypedates maculatus* (Gray, 1830) – are a specialized adaptation to reduce dehydration in arid environments (Amey and Grigg 1995, Lillywhite 2004, Barbeau and Lillywhite 2005, Gomez et al. 2006). These lipids are spread over the body by complex self-wiping behavior to form an

effective barrier that reduces evaporative water loss (Barbeau and Lillywhite 2005).

In this study, clusters of glands were observed in the ventrolateral integument of *O. albicans*, *O. angrensis*, *O. flavoguttata*, *S. hayii*. These glandular accumulations are probably present in a specialized region of the integument that provides special functions.

Clusters of tubuloalveolar alveoli in the ventral integument occurs in *Cycloramphus fuliginosus* (Gonçalves and de Brito-Gitirana 2008), being related to parental care. In *Rhinella icterica* (de Brito-Gitirana et al. 2007, Almeida et al. 2007) and in *R. ornata* (Felsemburgh et al. 2009), glandular aggregates constitute the parotoid gland, whose secretion is related to chemical defense against predators and parasites (Croce et al. 1973, Clarke 1997, Sakate and Lucas de Oliveira 2000).

In all hylids, examined in this study, melanophores occurred in the dorsal integument, but they did not always occur in the ventral integument. In contrast to melanophores, iridophores were visualized only in the dorsal region of *O. albicans*, *O. angrensis*, *O. humilis*, and *S. similis*, while they occurred only in the ventral integument of *S. albicans*.

In all examined species, at least in the dorsal region of the integument, the Eberth-Katschenko (EK) layer was visualized as an acellular layer that was restricted to a region between the spongy and compact dermis. Moreover, the EK-layer was usually continuous in the dorsal integument, showing its typical basophilic and alcianophilic stainings, which were due to the glycoconjugate content. In *R. icterica* and *L. catesbeianus*, the EK-layer contained both dermatan sulfate and calcium, and occurred as scattered aggregates throughout the spongy dermis (Pelli et al. 2007, 2010). These mineral consists of calcium phosphate deposits (Katchburian et al. 2001). Moreover, calcium of the EK-layer was more concentrated in the dorsal integument of male toads, but no significant difference was detected in the integument of females (Azevedo et al. 2005). Elkan (1968) suggested that the absence of the EK layer in some anuran species may be correlated with the fixative type and storage time. Nevertheless, in this study, all hylids were fixed in the same manner, using the same fixative, and the presence or absence of the EK-layer varied according to the specimen and integument region. In addition, the EK-layer was absent in the ventral region of the integument of the species of *Scinax*, probably being a genus-specific feature. For some authors (Katchburian et al. 2001, Mangione et al. 2011), the EK-layer might be a remnant of an ancestral dermal skeleton. Toledo and Jared (1993) suggested that the calcium located in the EK layer participates in hydric balance, affecting the hydric absorption and retention. On the other hand, Azevedo and et al. (2007) demonstrated that hyaluronic acid (HA) occurs in the spongy dermis, suggesting that the entire spongy dermis acts as a hydric reservoir since HA, an important component of connective tissue matrices, is involved in promoting matrix assembly, tissue hydration and viscosity of some fluids (Laurent et al. 1996). However, the functional significance of the presence of calcium in the EK-layer remains unclear.

In *O. angrensis*, *O. flavoguttata*, *O. perpusilla*, *O. trapicheroi*, *S. similis* and *S. x-signatus*, cutaneous elevations were evident in the ventral integument, and were separated by a network of grooves. Cutaneous elevations were also noted in the ventral integument of *Hyla arborea* (Linnaeus, 1758) (Goniakowska-Wi-

talinska and Kubiczek 1998). In *R. icterica* and in *Proceratophrys boiei* (Wied-Neuwied, 1824), *Proceratophrys laticeps* Izecksohn & Peixoto, 1981, *Proceratophrys appendiculata* (Günther, 1873) and *Odontophrynus americanus* (Duméril & Bibron, 1841), these elevations were separated by a network of grooves that probably acts as a distribution system of water from the ventral to dorsal surface of the integument (Azevedo and de Brito-Gitirana 2005, Felsemburgh et al. 2007). According to Lillywhite and Licht (1974), grooves can work as water distribution channels by a capillarity mechanism from one integument surface to another. Several authors have proposed that water distribution keeps the integument moist, protecting the animal against desiccation (Parakkal and Matoltsy 1964, Machin 1969, Duellman and Trueb 1994, de Brito-Gitirana and Azevedo 2005).

Although the usual patterns observed in the *Ololygon* and *Scinax* species, their integuments revealed histological characteristics. Thus, histological methods can be efficient to help characterize and differentiate of anuran integuments, thereby improving their taxonomy.

ACKNOWLEDGMENTS

We thank all colleagues for the help during fieldwork at Serra dos Órgãos National Park and the park staff for logistic support and permission for fieldwork. We also thank Conselho Nacional de Desenvolvimento Científico e Tecnológico (CNPq) and Fundação Carlos Chagas Filho de Amparo à Pesquisa do Estado do Rio de Janeiro (FAPERJ). TSS thanks support from CNPq (process 304374/2016-4).

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 Submitted: 4 May 2016

Received in revised form: 28 November 2016

Accepted: 22 December 2016

Editorial responsibility: Carolina Arruda Freire

Author Contributions: HMA, LBG and TSS participated equally in the preparation of this article.

Competing Interests: The authors have declared that no competing interests exist.